

SUPPLEMENTARY INFORMATION: The SORN is now obsolete and is being rescinded.

SYSTEM NAME AND NUMBER:

eLease. GSA/PBS-5.

HISTORY:

73 FR 22414, April 25, 2008.

Richard Speidel,

Chief Privacy Officer, Office of the Deputy Chief Information Officer, General Services Administration.

[FR Doc. 2023-10500 Filed 5-16-23; 8:45 am]

BILLING CODE 6820-34-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Office of Refugee Resettlement Unaccompanied Refugee Minors Program Application and Withdrawal of Application or Declination of Placement Form (OMB #0970-0550)

AGENCY: Office of Refugee Resettlement, Administration for Children and

Families (ACF), Department of Health and Human Services (HHS).

ACTION: Request for public comments.

SUMMARY: The Office of Refugee Resettlement (ORR) is requesting a 3-year extension with revisions of the Unaccompanied Refugee Minors (URM) Program Application and Withdrawal of Application or Declination of Placement Form (OMB #0970-0550, expiration 08/31/2023). Proposed revisions include additional instructions, a small number of new questions, dropping a few questions, and rephrasing existing questions.

DATES: *Comments due within 30 days of publication.* OMB must make a decision about the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication.

ADDRESSES: Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to www.reginfo.gov/public/do/

PRAMain. Find this particular information collection by selecting “Currently under 30-day Review—Open for Public Comments” or by using the search function. You can also obtain copies of the proposed collection of information by emailing infocollection@acf.hhs.gov. Identify all emailed requests by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: The URM Program Application is completed on behalf of unaccompanied children in the United States who are applying for entry into the URM Program. The application includes biographical data and information on the child’s needs to support placement efforts. The Withdrawal of Application or Declination of Placement Form is completed when a child is no longer interested in entering the URM Program or is not interested in entering the placement they were offered.

Respondents: Case managers, attorneys, or other representatives working with unaccompanied children who are eligible for the URM Program.

ANNUAL BURDEN ESTIMATES

| Instrument | Total number of respondents | Total number of responses per respondent | Average burden hours per response | Total burden hours | Annual burden hours |
|--|-----------------------------|--|-----------------------------------|--------------------|---------------------|
| Unaccompanied Refugee Minors Program Application | 450 | 3 | 1.5 | 2,025 | 675 |
| Withdrawal of Application or Declination of Placement Form | 50 | 3 | 0.2 | 30 | 10 |

Estimated Total Annual Burden Hours: 685.

Authority: 8 U.S.C. 1522(d).

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2023-10539 Filed 5-16-23; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2023-D-1103]

Compliance Policy Guide Sec. 555.250 Major Food Allergen Labeling and Cross-Contact; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of availability.

SUMMARY: The Food and Drug Administration (FDA or we) is announcing the availability of a draft Compliance Policy Guide entitled “Sec.

555.250 Major Food Allergen Labeling and Cross-contact.” The draft guidance, when finalized, will replace existing guidance for FDA staff on FDA’s enforcement policy regarding major food allergen labeling and cross-contact.

DATES: Submit either electronic or written comments on the draft guidance by July 17, 2023 to ensure that we consider your comment on the draft guidance before we begin work on the final version of the guidance.

ADDRESSES: You may submit comments on any guidance at any time as follows:

Electronic Submissions

Submit electronic comments in the following way:

- *Federal eRulemaking Portal:* <https://www.regulations.gov>. Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to <https://www.regulations.gov> will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your

comment does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else’s Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on <https://www.regulations.gov>.

- If you want to submit a comment with confidential information that you do not wish to be made available to the public, submit the comment as a written/paper submission and in the manner detailed (see “Written/Paper Submissions” and “Instructions”).

Written/Paper Submissions

Submit written/paper submissions as follows:

- *Mail/Hand Delivery/Courier (for written/paper submissions):* Dockets Management Staff (HFA-305), Food and